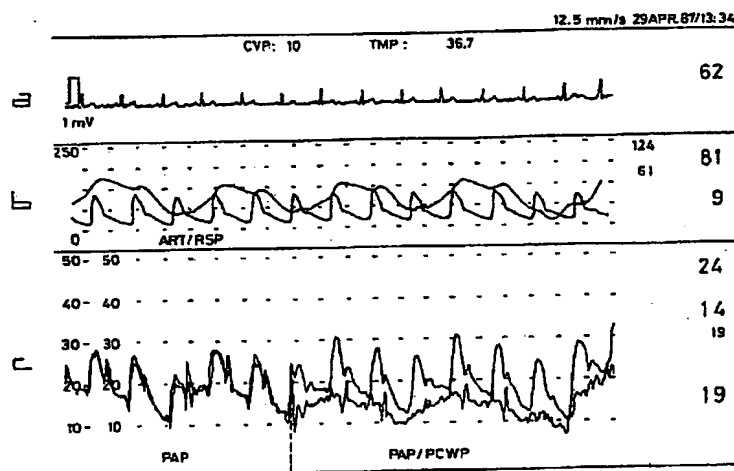




INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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(54) Title: METHOD FOR CARRYING OUT HEMODYNAMIC MEASUREMENTS ON A PATIENT AND FLOW-DIRECTED BALLOON CATHETER USED FOR THIS



(57) Abstract

A method for carrying out hemodynamic measurements on a patient, using a flow-directed balloon catheter (1) which is connected to a measuring unit and is at least provided with a distal measuring lumen (15) and a balloon-inflating lumen (16), comprising the steps of inserting the catheter by way of a suitable vein and further by way of the right atrium and ventricle into the pulmonary artery until the distal end with the inflated balloon in wedge position is stuck in a branch thereof and closes off the latter. A proximal measuring lumen (18) is provided a few centimetres apart from the distal measuring lumen (15). Then the pulmonary arterial pressure (PAP) is measured, as long as the catheter has not yet arrived in the wedge position, via both the proximal and distal measuring lumina and the two relevant, virtually identical pressure curves are simultaneously recorded. In the wedge position of the catheter with the balloon inflated, the pulmonary arterial pressure (PAP) and the pulmonary capillary wedge pressure (PCWP) via both said measuring lumina are measured respectively.

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AT	Austria	FR	France	ML	Mali
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Method for carrying out hemodynamic measurements on a patient and flow-directed balloon catheter used for this.

The invention relates to a method for carrying out hemodynamic measurements on a patient, using a flow-directed balloon catheter, which is connected to a control and pressure-measuring unit and is at least provided with a distal measuring lumen and a balloon-inflating lumen, comprising the steps of inserting the catheter by way of a suitable vein until the distal end is positioned inside the thorax, and of subsequently inflating the balloon and of inserting the catheter further by way of the right atrium and ventricle into the pulmonary artery until the distal end with the balloon in wedge position is stuck in a branch thereof and closes off the latter. By measuring lumen, the opening end of a through tube is meant, which tube is connected at the other end to the control and pressure-measuring unit.

This method and the catheter used for this are known, inter alia, from the article "Hemodynamic Monitoring" by N. Buchbinder and W. Ganz in "Anesthesiology", August 1976, vol 45, no. 2, pages 146-155.

In this known method and catheter, a number of hemodynamic measurements are carried out in order to gain an impression therewith of the functioning of heart and circulation. The pulmonary arterial pressure (PA pressure) and the pulmonary capillary wedge pressure (PCW pressure) play a very important role here. The pulmonary capillary wedge pressure is regarded as being a good measure of the end-diastolic pressure in the left ventricle because there is a continuous head of fluid (blood) between the left atrium and the catheter end in wedge position. As a result the PCW pressure can be designated as the filling pressure in the left atrium. The functioning of the left ventricle as a pump is determined to a considerable extent by the end-diastolic volume in

the left ventricle. For a satisfactory approximation of this volume, the end-diastolic pressure in this left ventricle can be used. Catheterization of an artery would be necessary to measure this pressure directly. Instead of this, the PCW pressure, which
5 can be assessed by routine measurement with hemodynamic monitors, is used. It is evident that a partial or unsatisfactorily controlled wedge position of the catheter makes the measurement unreliable. A partial wedge position will generally be recognisable by the pressure curve shown on a monitor. However, this is not always the
10 case because PCW pressure curves often do not show the "ideal picture" and, because there is great variability in PCW pressure curves.

Additionally, the known method of carrying out hemodynamic measurements has the problem that the PA pressure is measured during
15 introduction of the catheter with the inflated balloon into the pulmonary artery and that the PCW pressure is measured at a subsequent instant when the wedge position of the catheter is reached. It also holds that the PA pressure is measured when the catheter has been brought into position without the balloon being
20 inflated and that the PCW pressure is measured at a subsequent instant, after inflating the balloon. Due to the fact that these curves are recorded only at consecutive instants in this way they can therefore be assessed and evaluated on the monitor exclusively with consideration of the time difference. A further problem which
25 occurs is that a catheter with non-inflated balloon may come in a wedge position which is displaced upwards, which may not be noticed or is not directly noticeable because the PCW pressure curve cannot always satisfactorily be distinguished from the PA pressure curve which is measured first. Such an upward displaced wedge position,
30 which may be caused with a non-inflated balloon by some artefact or other, may have very adverse effects on the blood through-flow in the pulmonary artery of the patient.

The invention aims at overcoming the above-mentioned problems and at providing an exceptionally efficient and safe method
35 and a catheter used for this, by which the safety of insertion and positioning of the catheter and the quality and control of the pulmonary pressure measurement are guaranteed.

This is achieved, according to the invention, in a method of the type mentioned at the outset in a manner such that a proximal measuring lumen is provided in the catheter a few centimetres apart from the distal measuring lumen, and that during the insertion of the catheter into the pulmonary artery, as long as the catheter has not yet arrived in the wedge position, the pulmonary arterial pressure is measured both via the proximal measuring lumen and via the distal measuring lumen and that the two relevant pressure curves which are virtually identical are simultaneously recorded by the measuring unit and/or represented thereon.

In a further embodiment of the method according to the invention, the method comprises the step of subsequently measuring, in the wedge position of the catheter with the balloon inflated, the pulmonary arterial pressure and the pulmonary capillary wedge pressure via the proximal measuring lumen and via the distal measuring lumen respectively and of simultaneously recording the two relevant, mutually different pressure curves with the measuring unit and/or representing them thereon. By this means, the moment at which the catheter arrives in the wedge position can be clearly detected by the two said curves diverging, as a result of which the simultaneous measurement of PA pressure and PCW pressure relative to each other is independent of interfering cardiac and respiratory influences.

In another embodiment of the method according to the invention, the method comprises the step of incidentally inflating the balloon in the wedge position of the catheter, the pressures measured simultaneously via the two measuring lumina in the uninflated and inflated state of the balloon producing pressure curves, which are virtually identical and mutually different, respectively, on the measuring unit as a result of which the wedge position is exactly determined and a reliable measurement of the two said pressures is guaranteed. Thereafter, in a spontaneously assumed and upward-displaced wedge position of the catheter with the balloon uninflated, the pulmonary arterial pressure and pulmonary capillary wedge pressure measured simultaneously via the proximal measuring lumen and via the distal measuring lumen will produce different pressure curves on the measuring unit as a result of which

the moment at which the catheter arrives spontaneously in an upward-displaced wedge position with the balloon uninflated can be clearly detected by the divergence of the two said curves and an alarm can be given.

5 In yet another embodiment of the method according to the invention, the method comprises the step of measuring, during the insertion of the catheter by way of the right atrium and ventricle into the pulmonary artery or during removal of the catheter therefrom, the pressure on one side and on the other side of the
10 tricuspid valve and/or the pulmonary valve, by means of the proximal and distal measuring lumina, so that the pressure gradient over the said valve(s) is recorded and/or represented on the measuring unit by means of two pressure curves.

The flow-directed balloon catheter mentioned in the outset
15 is designed for carrying out the abovementioned methods in a manner such that a proximal measuring lumen is provided a few centimetres apart from the distal measuring lumen which makes it possible to measure the pulmonary arterial pressure and the pulmonary capillary wedge pressure simultaneously in the wedge position of the
20 catheter. Present catheter may also be implemented such that each measuring lumen includes a pressure transducer, f.i. in chip form. Said transducer directly converts the pressure, measured at said lumen, into an electrical signal. Said signal is transmitted to the measuring unit instead that the pressure, otherwise present in the
25 lumen and corresponding tube, is converted into a signal via a transducer in the measuring unit per se.

It is known in practice to provide such a flow-directed balloon catheter with a proximal lumen for measuring pressures, this being provided, however, at a distance of 25 to 30 cm from the
30 distal tip. This proximal lumen is solely intended for measuring the right atrial pressure or the central venous pressure when the distal tip of the catheter is located in the pulmonary artery. Furthermore, particular fluids for infusion or medicines may be administered via this lumen.

35 The invention will be explained in detail by means of an embodiment with reference to the drawings, in which:

Figure 1 shows a simplified diagram of the systemic blood

circulation in human beings;

Figure 2 shows a view of a known flow-directed balloon catheter;

5 Figure 3 shows a curve exemplifying the blood pressure measured via the catheter from the right atrium to the wedge position;

Figure 4 shows a view of the flow-directed catheter according to the invention;

10 Figure 5 shows curves of the pulmonary arterial pressure and the pulmonary capillary wedge pressure measured according to the state of the art and of some other parameters;

Figure 6 shows curves of the pulmonary arterial pressure and the pulmonary capillary wedge pressure measured according to the invention, and of some other parameters;

15 Figures 7 and 8 show curves of the pressures measured according to the invention on either side, of the tricuspid valve and the pulmonary valve, and of some other parameters; and

Figures 9 and 10 show a few other curves measured according to the invention.

20 Figure 1 shows diagrammatically the systemic blood circulation with the heart in diastole, i.e. with the ventricles 7 and 9 relaxed, the pulmonary valve 10 and aortic valve 11 closed and the tricuspid valve 12 and mitral valve 13 opened. In this, 1 designates a catheter inserted into the pulmonary artery 3. Part of
25 the pulmonary circulation, the systemic circulation of the body, the right atrium, the right ventricle, the left atrium, the left ventricle and the pulmonary vein are indicated diagrammatically by 4, 5, 6, 7, 8, 9 and 14, respectively. It can be seen how the flow-directed balloon catheter 1 with the balloon 2 inflated has
30 arrived near the wedge position in the pulmonary artery 3.

Figure 2 shows a known catheter provided with a distal measuring lumen 15, a balloon-inflating lumen 16 and a thermistor 17 (the function of which is explained hereinafter). During insertion
35 of the catheter the distal measuring lumen can be used to measure the pressure consecutively in the right atrium, right ventricle and thereafter in the pulmonary artery as is shown in Figure 3. This figure shows at the bottom specifically, viewed from the left, in

succession the right atrium pressure (RAP), right ventricular pressure (RVP), pulmonary arterial pressure (PAP) and pulmonary capillary wedge pressure (PCWP) during insertion of the catheter and the pulmonary arterial pressure (PAP) again during removal of the catheter or deflation of the balloon. This figure furthermore shows the electrocardiogram (EKG), the radial arterial pressure (ART) and respiration (RSP) of said patient.

Figure 4 shows the catheter according to the invention which is additionally provided with a proximal measuring lumen 18 a few centimetres apart from the distal measuring lumen 15. As indicated before each lumen may include a pressure (chip) transducer. This embodiment makes it possible to measure simultaneously the PA pressure and PCW pressure and the pressures at either side of the tricuspid valve and on either side of the pulmonary valve as indicated by Figures 5 to 8.

As mentioned before, the position of the distal end of the catheter may represent a possible risk for the patient. In fact, the catheter may spontaneously move into an upward-displaced wedge position with the balloon uninflated. As a result, the blood supply to the region of the lung located behind this is shut off, and the consequence may be a pulmonary infarct. Such an unexpected displacement of the catheter tip without inflated balloon into an upward-displaced wedge position is not always noticed or it is not always directly recognisable from the PA pressure curve or the PCW pressure curve which then occurs. However, in this situation the new catheter can contribute to an alarm being given by the control and pressure measuring unit.

Figure 5 consecutively shows the electrocardiogram (Figure 5a), the radial arterial pressure (ART) and respiration (RSP) of a patient (Figure 5b), and the pressure measured in the pulmonary artery (Figure 5c) in mm Hg. This PA pressure measured with the known catheter passes over to the PCW pressure during further insertion of the catheter with inflated balloon or when the catheter is already in the wedge position after inflation of the balloon. CVP and TMP means central venous pressure and temperature respectively.

Figure 6 consecutively shows the electrocardiogram (Figure 6a), the radial arterial pressure (ART) and respiration (RSP) of a

patient (Figure 6b), and the pressures measured by the distal and proximal measuring lumina of the catheter according to the invention (Figure 6c) in the pulmonary artery. During further insertion of the catheter with the balloon inflated or when the catheter is already in the wedge position after inflation of the balloon, the two curves will diverge because the distal measuring lumen will measure the PCW pressure and the proximal measuring lumen will measure the PA pressure.

Due to the fact that the PA and PCW pressure curves can be measured simultaneously, control of the pulmonary pressure measurement is considerably improved. When the catheter is in position, but not in the wedge (balloon not inflated), the PA pressure curve and the PCW pressure curve converge. If this is not the case the catheter has come spontaneously in an upward-displaced wedge position or the two pressure systems are no longer exactly synchronous (artefact transducer, incorrect calibration procedure, etc.). In the case of a single pulmonary pressure curve it will not be possible to detect these deviations as quickly.

Figure 7 consecutively shows for a patient: the electrocardiogram (Figure 7a), the respiration (Figure 7b) and the simultaneously measured right atrial pressure (RAP: bottom curve) and right ventricular pressure (RVP: top curve) (Figure 7c). The two curves in Figure 7c are measured by the proximal measuring lumen and distal measuring lumen of the catheter according to the invention.

Figure 8 consecutively shows for a patient: the electrocardiogram (Figure 8a), the respiration (Figure 8b) and the simultaneously measured right ventricular pressure (RVP: bottom curve) and pulmonary pressure (PAP: top curve) (Figure 8c). The two curves in Figure 8c are measured by the proximal measuring lumen and distal measuring lumen of the catheter according to the invention.

As indicated in Figures 7 and 8, the pressure on one side and on the other side of the tricuspid valve and the pulmonary valve can be measured and recorded simultaneously via the proximal and distal measuring lumina during insertion or removal of the catheter. This makes it possible to diagnose stenosis or valvular insufficiency.

The good quality of the pulmonary pressure measurement described above moreover results in the following.

As has been illustrated, the measurement becomes independent of interfering cardiac and respiratory influences due to the simultaneous recording of the PAP curve and the PCWP curve. The intrathoracic pressure changes and hemodynamic fluctuations are completely parallel in the two curves. As an example reference is made to Figure 9 which shows an interfering cardiac influence (Figure 9a), i.e. a ventricular extrasystole, and an interfering respiratory influence (Figure 9b), while said PAP and PCWP curves (Figure 9c) relative to each other are nevertheless satisfactorily interpretable.

This is also of great importance for calculating the pulmonary vascular resistance (PVR). When measurement of the PAP and PCWP is not carried out in an exact and synchronous manner this may incorrectly result in a negative deflection of the PVR. Unphysiological data may be quickly obtained especially when relying exclusively on digital reading on the monitor and when the fundamental curves are not used for quality control.

PVR is determined using the equation $PVR = (\overline{PAP} - \overline{LAP}) / CO$, wherein LAP is the left atrial pressure and CO the cardiac output. This cardiac output CO is measured via the known thermodilution method, and for this the catheter is provided with a thermistor. The left atrial pressure (LAP) can be replaced, as indicated in the introduction, approximately by the pulmonary capillary wedge pressure (PCWP).

In the context of analyzing the pulmonary vascular resistance, determined by a viscosity factor to be measured and a vascular factor not to be measured, correct measurement of PCWP and calculation of PVR is of fundamental value for specifying the medical indication. Reduction of iatrogenic complications is a further safety aspect which is not unimportant.

If the PAP curve and the PCWP curve are subtracted from each other as shown in Figure 10d, a " $\Delta(PAP-PCWP)$ " curve can be seen which is completely independent of the respiration (Figure 10b) and therefore independent of the intrathoracic pressure changes (Figure 10c) due to respiration, and which is representative of the cardiac

output and the pulmonary vascular resistance ($CO \times PVR$).

5 One of the reasons why pulmonary pressures are difficult to read is that the distal tip of the catheter sometimes "sways" ("catheter-whip effect") and that no satisfactory PAP curve can be distinguished. When the balloon is inflated with preservation of the PAP and PCWP measurements, the distal tip of the catheter is fixed and this problem is largely solved. The possibility of PAP measurement with inflated balloon is lost in the classical catheter.

CLAIMS

1. Method for carrying out hemodynamic measurements on a patient, using a flow-directed balloon catheter, which is connected to a control and pressure-measuring unit and is at least provided with a distal measuring lumen and a balloon-inflating lumen,
5 comprising the steps of inserting the catheter by way of a suitable vein until the distal end is positioned inside the thorax and of subsequently inflating the balloon, and of inserting the catheter further by way of the right atrium and ventricle into the pulmonary artery until the distal end with the inflated balloon in wedge
10 position is stuck in a branch thereof and closes off the latter, a proximal measuring lumen being provided a few centimetres apart from the distal measuring lumen, the pulmonary arterial pressure (PAP) being measured, during insertion of the catheter into the pulmonary artery as long as the catheter has not yet arrived in the wedge
15 position, via both the proximal measuring lumen and the distal measuring lumen and the two relevant, virtually identical pressure curves being simultaneously recorded by the measuring unit and/or represented thereon.
2. Method according to claim 1, comprising the step of
20 measuring, in the wedge position of the catheter with the balloon inflated, the pulmonary arterial pressure (PAP) and the pulmonary capillary wedge pressure (PCWP) via the proximal measuring lumen and via the distal measuring lumen, respectively, and of simultaneously recording the two relevant, mutually different pressure curves with
25 the measuring unit and/or representing them thereon.
3. Method according to claim 2, comprising the step of incidentally inflating the balloon in the wedge position of the catheter, the pressures measured simultaneously via the two
30 measuring lumina in the uninflated and inflated state of the balloon producing virtually identical and mutually different pressure curves on the measuring unit as a result of which the wedge position is exactly determined and a reliable measurement of the two said pressures is guaranteed.
4. Method according to claim 3, comprising the step of
35 simultaneously measuring the pulmonary arterial pressure (PAP) and

pulmonary capillary wedge pressure (PCWP) via the proximal measuring lumen and via the distal measuring lumen in a spontaneously assumed and upward-displaced wedge position of the catheter with the uninflated balloon producing thereby mutually different pressure curves on the measuring unit, as a result of which the moment at which the catheter without inflated balloon spontaneously arrives in an upward-displaced wedge position can be clearly detected due to the two said curves diverging and an alarm can be given.

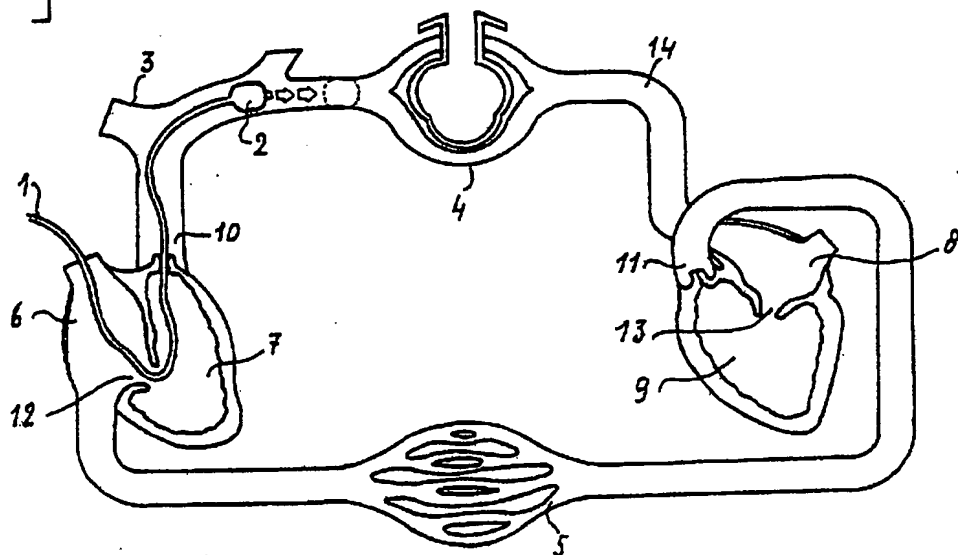
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15
5. Method according to claim 1, comprising the step of measuring, during the insertion of the catheter by way of the right atrium and ventricle into the pulmonary artery or during removal of the catheter therefrom, the pressure on one side and on the other side of the tricuspid valve and/or the pulmonary valve by means of the proximal and distal measuring lumina, so that the pressure gradient over the said valve(s) is recorded and/or represented on the measuring unit by means of two pressure curves.

20
6. Method according to claim 2 or 3, in which the catheter has a thermistor lumen near the distal end for a thermodilution measurement in order to determine the cardiac output (CO), the pulmonary vascular resistance $PVR = (PAP - LAP)/CO$ being determined (wherein LAP is the left atrial pressure) by $LAP \sim PCWP$, the intra-thoracic pressure changes, hemodynamic and respiratory fluctuations which may occur during the measurement of PAP and PCWP having no influence.

25
30
7. Flow-directed balloon catheter for hemodynamic measurements for use with the method according to one of the preceding claims, in which the catheter is at least provided with a distal measuring lumen and a balloon-inflating lumen, characterized in that a proximal measuring lumen is provided a few centimetres apart from the distal measuring lumen which makes it possible to measure the pulmonary arterial pressure (PAP) and the pulmonary capillary wedge pressure (PCWP) simultaneously in the wedge position of the catheter.

35
8. Catheter according to claim 7, in which each measuring lumen comprises a pressure (chip) transducer.

fig-1

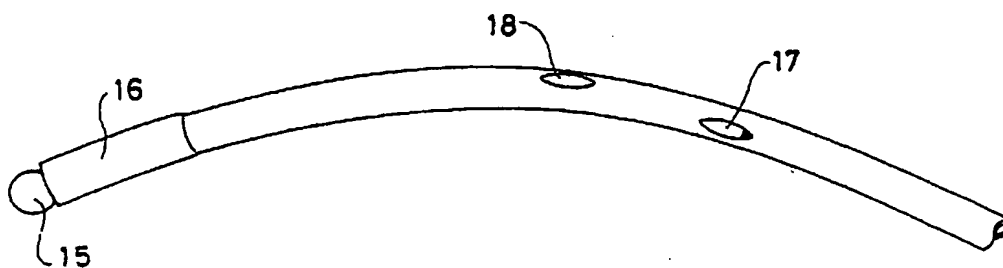


2/9

fig-2

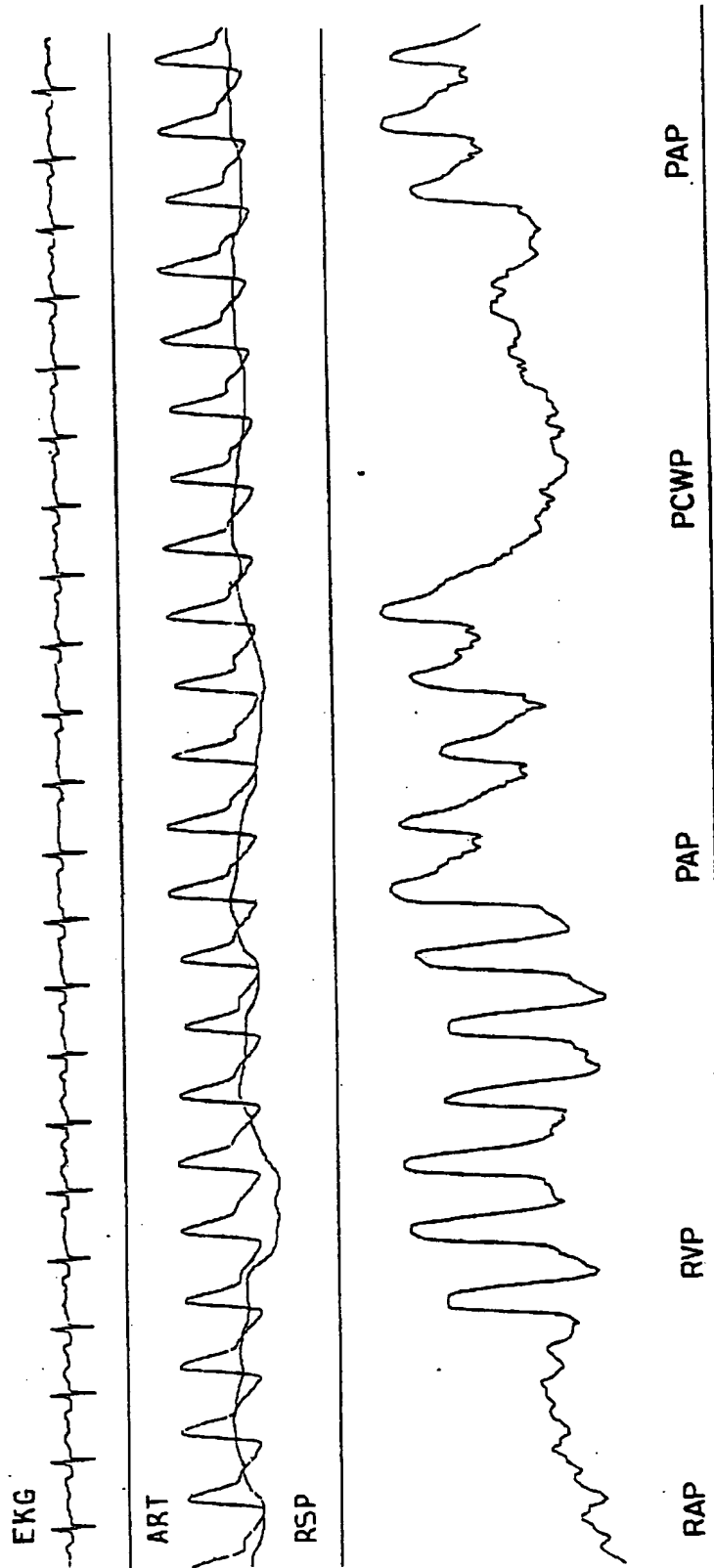


fig-4



3/9

fig-3



4/9

fig-5

12.5 mm/s 29APR.87/13.31

CVP: 9 TMP: 36.8

57 beats/min.

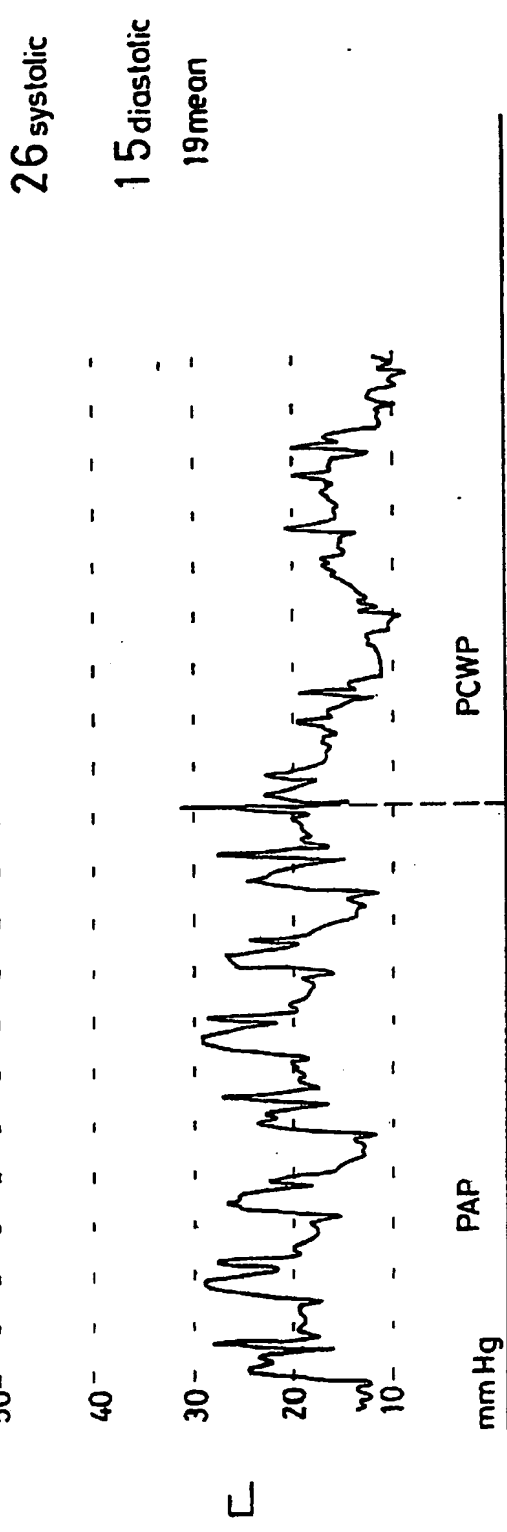
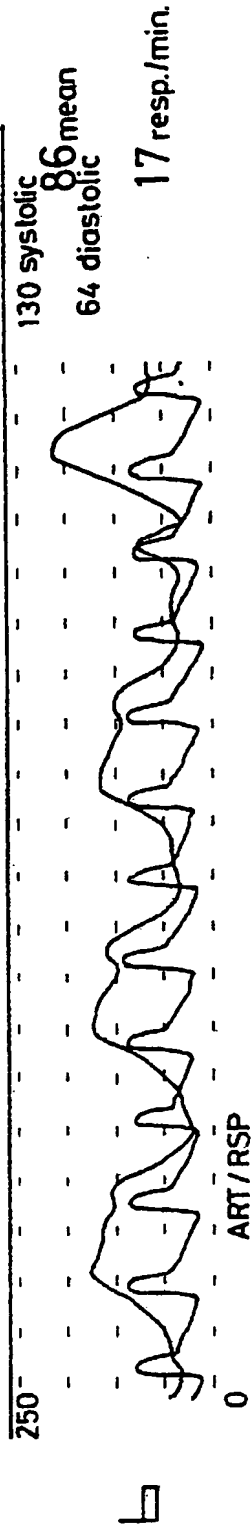
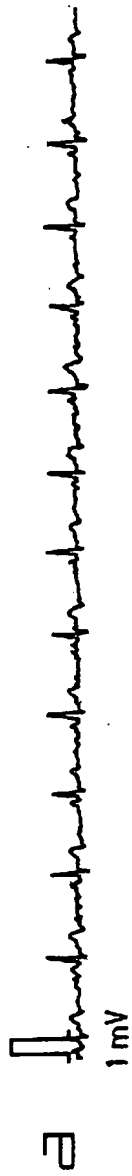
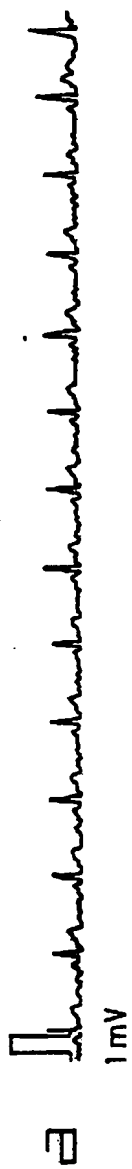


fig-6

12.5 mm/s 29APR 87/13:34

CVP: 10 TMP: 36.7

62



250 124 81 61

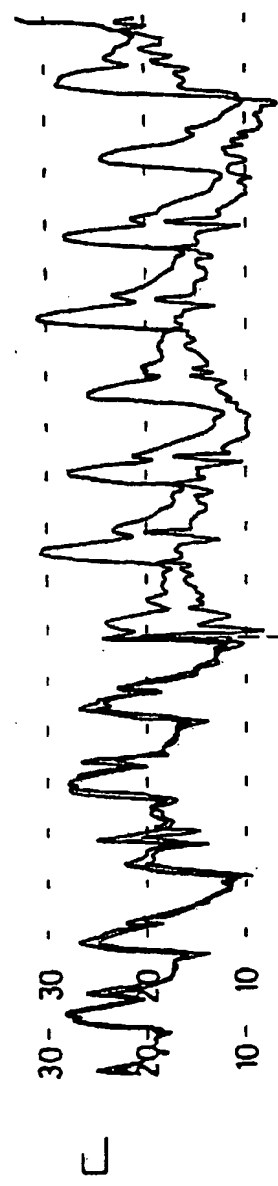


24

14

19

19



PAP/PCWP

PAP

fig-7

24 APR 87/11:43

CVP: 8 TMP: 36.2

65



18



RSP

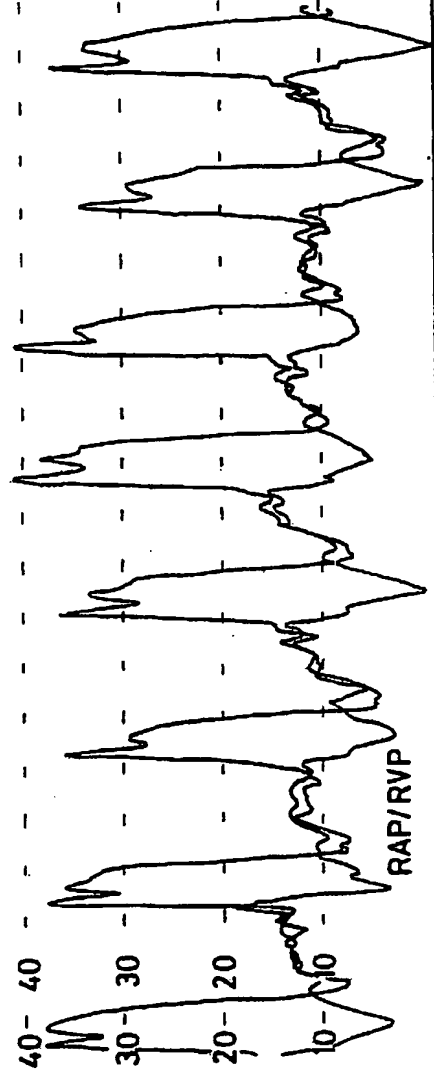
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6

18

8



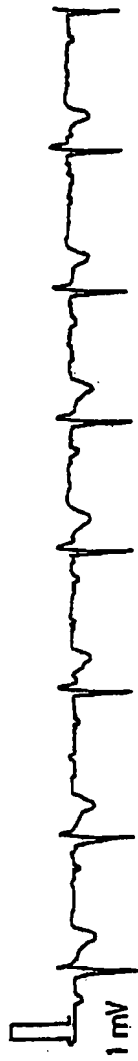
7/9

fig - 6

24 APR 87/11:42

CVP: 53 TMP: 36.2

69



18



31



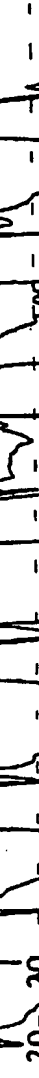
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18



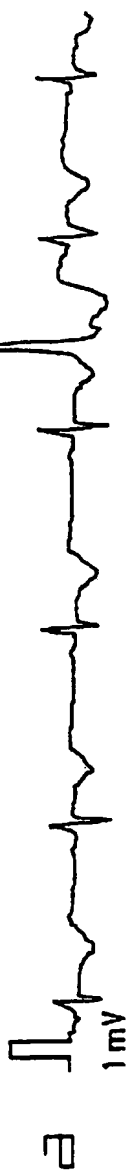
8/9

fig-9

06FEB.87/08:57

CVP: 15 TMP: 36.9

51

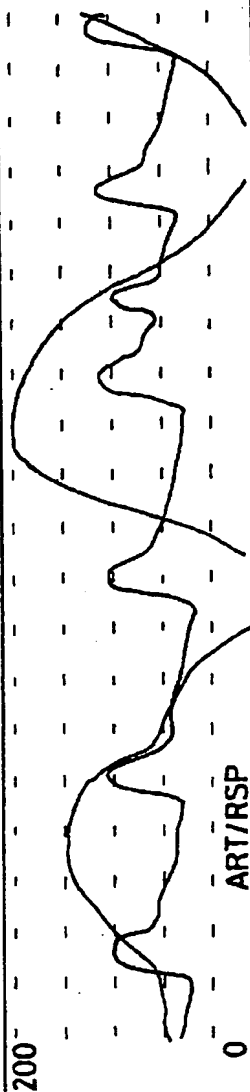


131

86

64

21



ART/RSP

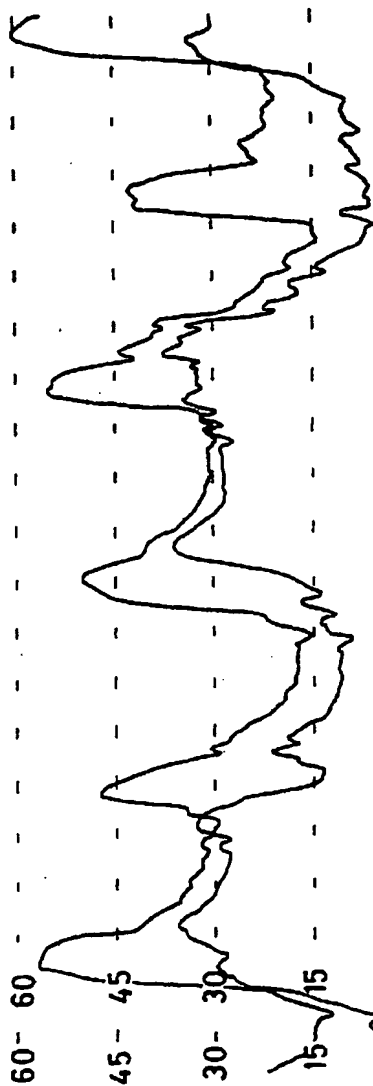
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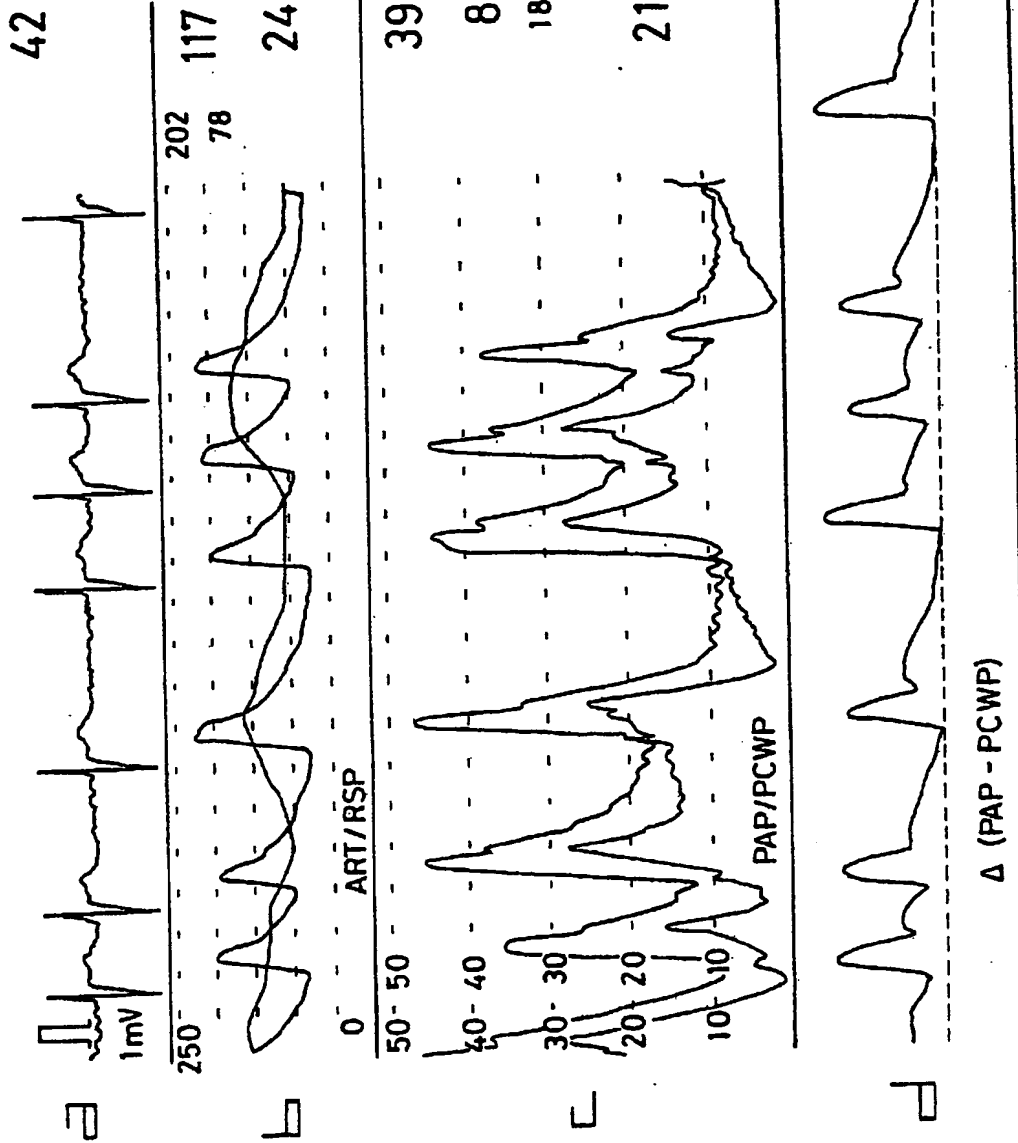
PAP/PCWP

9/9

Fig-10

18 JUN.87/20:57

CVP: 5 TMP: 39.7



INTERNATIONAL SEARCH REPORT

International Application No PCT/NL 88/00031

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) *

According to International Patent Classification (IPC) or to both National Classification and IPC

IPC⁴: A 61 B 5/02

II. FIELDS SEARCHED

Minimum Documentation Searched *

Classification System |

Classification Symbols

IPC⁴

A 61 B; A 61 M

Documentation Searched other than Minimum Documentation
to the extent that such Documents are included in the Fields Searched *

III. DOCUMENTS CONSIDERED TO BE RELEVANT*

Category *	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
Y	US, A, 3995623 (L.W. BLAKE et al.) 7 December 1976, see the whole document	1,2,5,7
A	--	3,4,6
Y	US, A, 4543961 (D.C. BROWN) 1 October 1985, see abstract; column 1, lines 20-40	1,2,5,7
A	--	4,8
A	Proceedings of the IEEE, vol. 65, no. 5, May 1977, (New York, US) J.P. Murgo et al.: "Physiologic signal acquisition and processing for human hemodynamic research in a clinical cardiac-catheterization laboratory", pages 696-702, see page 692, paragraph "II. Signal Acquisition"; figures 1-3	1-8

* Special categories of cited documents: ¹⁰

"A" document defining the general state of the art which is not
considered to be of particular relevance

"E" earlier document but published on or after the international
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"T" later document published after the international filing date
or priority date and not in conflict with the application but
cited to understand the principle or theory underlying the
invention

"X" document of particular relevance; the claimed invention
cannot be considered novel or cannot be considered to
involve an inventive step

"Y" document of particular relevance; the claimed invention
cannot be considered to involve an inventive step when the
document is combined with one or more other such docu-
ments, such combination being obvious to a person skilled
in the art.

"&" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search

21st September 1988

Date of Mailing of this International Search Report

07 OCT 1988

International Searching Authority

EUROPEAN PATENT OFFICE

Signature of Authorized Officer

 P.C.G. VAN DER PUTTEN

FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

- | | | |
|---|--|-------|
| A | IEEE Transactions on Sonics and Ultrasonics, vol. SU-27, no. 6, November 1980 (New York, US) R.W. Martin et al.: "An ultrasonic catheter for intra-vascular measurement of blood flow: technical details", pages 277-281, see page 277, paragraph "Catheterization requirements"; figure 1 | 1,3,7 |
| A | US, A, 4554927 (T.J. FUSSELL) 26 November 1985, see abstract; column 4, lines 22-64; figures 1-5 | 8 |

V. ☒ OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND incompletely searchable

This international search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1. ☒ Claim numbers * because they relate to subject matter not required to be searched by this Authority, namely:

* 1-6

See PCT Rule 39.1(iv): Methods for treatment of the human or animal body by surgery or therapy, as well as diagnostic methods.

2. ☐ Claim numbers because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. ☐ Claim numbers because they are dependent claims and are not drafted in accordance with the second and third sentences of PCT Rule 6.4(a).

VI. ☐ OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING ²

This International Searching Authority found multiple inventions in this international application as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application.
2. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims:
3. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:
4. ☐ As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

Remark on Protest

- ☐ The additional search fees were accompanied by applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.

NL 8800031
SA 22914

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report.
The members are as contained in the European Patent Office EDP file on 04/10/88
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US-A- 3995623	07-12-76	None	
US-A- 4543961	01-10-85	None	
US-A- 4554927	26-11-85	None	

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For more details about this annex : see Official Journal of the European Patent Office, No. 12/82